Application No. PC1 03/01595 Attorney Docket No. PG4715USW

## AMENDMENT OF THE CLAIMS:

- 1. (Currently amended) A dry powder pharmaceutical composition for inhalation therapy comprising salmeterol or a pharmaceutically acceptable salt thereof and fluticasone propionate, an excipient and a derivatised derivatized carbohydrate in particulate form.
- 2. (Currently amended) A dry powder pharmaceutical composition according to claim 1 in which salmeterol is present as its 1-hydroxy-2-naphthoate (xinofoate)-salt.
- 3. (Currently amended) A dry powder pharmaceutical composition according to claim 1 or 2 in which the derivatised derivatized carbohydrate is a mono or disaccharide in which at least one hydroxyl group of the carbohydrate group is substituted with a hydrophobic moiety via either ester or ethers linkages.
- 4. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1 3 claim 1 in which the derivatised derivatized carbohydrate is a carbohydrate selected from fructose, glucose, mannitol, maltose, mannitol, trehalose, cellobiose, lactose and sucrose in which at least one hydroxyl group of said carbohydrate is substituted by a straight or branched hydrocarbon chain comprising up to 20 carbon atoms.
- 5. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1—4 claim 1 in which the derivatised derivatized carbohydrate is selected from the group consisting of cellobiose octaacetate, sucrose octaacetate, glucose pentacetate, mannitol hexaacetate and trehalose octaacetate.
- 6. (Currently amended) A dry powder pharmaceutical composition according to claim 1 in which the derivatised derivatized carbohydrate is α-D cellobiose octaacetate.

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- 7. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1 5 claim 1 in which the derivatised derivatized carbohydrate is present at a concentration of less than 10% of the total composition.
- 8. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1.—7 claim 1 in which the derivatised derivatized carbohydrate has an aerodynamic size in the range 1 20 μm.
- 9. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1—8 claim 1 in which one component of the excipient that has a particle size of less than 15µm (the fine excipient component) and another component of the excipient that has a particle size of greater than 20µm but lower than 150µm (the coarse excipient component).
- 10. (Original) A dry powder pharmaceutical composition according to claim 9 in which the fine and coarse excipient components are both lactose.
- 11. (Currently amended) A dry powder pharmaceutical composition according to any one of claims 1—10 claim 1 for use in therapy.
- 12. (Currently amended) A method of treatment or prophylaxis of respiratory disorders which comprises administering to a patient in need thereof a dry powder pharmaceutical composition according to any one of claims 1—10 claim 1.
- 13. (Cancelled) Use of a dry powder pharmaceutical composition according to any one of claims 1—10 in the manufacture of a medicament for the treatment of respiratory disorders.
- 14. (Currently amended) An inhalation device containing therein a dry powder pharmaceutical composition according to any one of claims 1—10 claim 1.
- 15. (Original) An inhalation device according to claim 14 in which the dry powder pharmaceutical composition is released from a pre-metered unit medicament pack.

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16. (Currently amended) A medicament pack for use in an inhalation device which comprises an elongate strip formed from a base sheet having a plurality of recesses spaced along its length and a lid sheet hermetically but peelably sealed thereto to define a plurality of containers, each container having therein an inhalable composition according to any one of claims 1—10 claim 1.

- 17. (Original) A medicament pack according to claim 16 wherein the strip is sufficiently flexible to be wound into a roll.
- 18. (Original) A medicament pack according to claim 16 wherein the lid sheet and base sheet have leading end portions which are not sealed to one another.
- 19. (Original) A medicament pack according to claim 18 wherein at least one of the said leading end portions is constructed to be attached to a winding means.
- 20. (Original) A medicament pack according to claim 16 wherein the hermetic seal between the base and lid sheets extends over their whole width.
- 21. (Original) A medicament pack according to claim 16 wherein the lid sheet may be peeled from the base sheet in a longitudinal direction from a first end of the said base sheet.
- 22. (Currently amended) A method of improving stability performance The use of particulate derivatised carbohydrates in dry powder pharmaceutical compositions comprising salmeterol or a pharmaceutically acceptable salt thereof and fluticasone propionate, said method including the step of including in said composition a particulate derivatized carbohydrate in order to improve stability performance.
- 23. (Currently amended)

  A method of eliminating or reducing the detrimental effect on fine particle dose experienced during storage of a dry powder pharmaceutical composition comprising salmeterol or a pharmaceutically acceptable salt thereof and fluticasone propionate, wherein said method comprises the step of including a The use of particulate derivatised derivatized earbohydrates carbohydrate

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in <u>said</u> dry powder pharmaceutical compositions <del>comprising salmeterol or a</del> <del>pharmaceutically acceptable salt there of and fluticasone propionate in order to eliminate or reduce the detrimental effect on fine particle dose caused on storage of said compositions.</del>

- 24. (Currently amended) The use according to method of claim 22 or 23 in which the particulate derivatized derivatized carbohydrate is cellobiose octaacetate.
- 25. (New) The method of claim 23 in which the particulate derivatized carbohydrate is cellobiose octaacetate.